Complete Summary

GUIDELINE TITLE

ACC/AHA guideline update on perioperative cardiovascular evaluation for noncardiac surgery: a report of the American College of Cardiology/American Heart Association Task Force on Practice Guidelines (Committee to Update the 1996 Guidelines on Perioperative Cardiovascular Evaluation for Noncardiac Surgery).

BIBLIOGRAPHIC SOURCE(S)

American College of Cardiology Foundation (ACCF), American Heart Association (AHA). ACC/AHA guideline update on perioperative cardiovascular evaluation for noncardiac surgery. A report of the American College of Cardiology/American Heart Association Task Force on practice guidelines (Committee to Update the 1996 Guidelines). Bethesda (MD): American College of Cardiology Foundation; 2002. 58 p. [390 references]

COMPLETE SUMMARY CONTENT

SCOPE

METHODOLOGY - including Rating Scheme and Cost Analysis RECOMMENDATIONS

EVIDENCE SUPPORTING THE RECOMMENDATIONS

BENEFITS/HARMS OF IMPLEMENTING THE GUIDELINE RECOMMENDATIONS QUALIFYING STATEMENTS

IMPLEMENTATION OF THE GUIDELINE

INSTITUTE OF MEDICINE (IOM) NATIONAL HEALTHCARE QUALITY REPORT CATEGORIES

IDENTIFYING INFORMATION AND AVAILABILITY

SCOPE

DISEASE/CONDITION(S)

- Cardiovascular diseases, including:
 - Coronary artery disease
 - Myocardial infarction
 - Angina pectoris
 - Heart failure
 - Arrhythmias and conduction defects
 - Hypertension
 - Cardiomyopathy
 - Valvular heart disease
 - Pulmonary vascular disease
- Presence of implanted pacemakers and implantable cardioverter defibrillators

GUIDELINE CATEGORY

Evaluation Management Risk Assessment

CLINICAL SPECIALTY

Anesthesiology
Cardiology
Emergency Medicine
Family Practice
Internal Medicine
Nuclear Medicine
Surgery

INTENDED USERS

Physicians

GUIDELINE OBJECTIVE(S)

- To provide a framework for considering cardiac risk of noncardiac surgery in a variety of patients and surgical situations
- To guide preoperative evaluation to determine the patient's current medical status, make recommendations concerning the evaluation, management and risk of cardiac problems over the entire perioperative period, and provide a clinical risk profile that the patient, his or her primary physician, anesthesiologist, and surgeon can use in making treatment decisions that may influence short- and long-term cardiac outcomes

TARGET POPULATION

Patients undergoing noncardiac surgery

INTERVENTIONS AND PRACTICES CONSIDERED

Risk Assessment/Prognosis

- 1. Clinical history
- 2. Physical examination
- 3. Stepwise approach to perioperative cardiac assessment (clinical markers, prior coronary evaluation and treatment, functional capacity, and surgery-specific risk)
- 4. Supplemental preoperative evaluation:
 - a. Resting left ventricular function
 - b. 12-lead electrocardiogram
 - c. Exercise or pharmacological stress testing
 - d. Myocardial perfusion imaging
 - e. Dobutamine stress echocardiography
 - f. Ambulatory electrocardiogram monitoring

g. Coronary angiography

Management

- 1. Perioperative therapy
 - a. Surgical coronary revascularization: preoperative coronary artery bypass grafting (CABG); percutaneous transluminal coronary angioplasty (PTCA)
 - b. Pharmacologic management: beta-blockers; alpha-2 agonists
- 2. Management of specific preoperative cardiovascular conditions
- 3. Anesthetic considerations and intraoperative management, including choice of anesthetic technique and agent, perioperative pain management, use of intraoperative nitroglycerin, transesophageal echocardiography, maintaining body temperature, and use of intra-aortic balloon counterpulsation devices
- 4. Perioperative surveillance, including use of intraoperative pulmonary catheters, and ST-segment monitoring, and surveillance for perioperative myocardial infraction
- 5. Postoperative and long-term management

MAJOR OUTCOMES CONSIDERED

- Positive and negative predictive value of tests for myocardial infarction or death
- Short- and long term cardiac outcomes, such as perioperative cardiovascular morbidity (e.g., myocardial infarction) and mortality (e.g., cardiac death)

METHODOLOGY

METHODS USED TO COLLECT/SELECT EVIDENCE

Hand-searches of Published Literature (Primary Sources) Hand-searches of Published Literature (Secondary Sources) Searches of Electronic Databases

DESCRIPTION OF METHODS USED TO COLLECT/SELECT THE EVIDENCE

The American College of Cardiology/American Heart Association Committee to Update the 1996 Guidelines on Perioperative Cardiovascular Evaluation for Noncardiac Surgery conducted a comprehensive review of the literature relevant to perioperative cardiac evaluation since the last publication of these guidelines in 1996. Literature searches were conducted in the following databases: PubMed/MEDLINE, EMBASE, the Cochrane Library (including the Cochrane Database of Systematic Reviews and the Cochrane Controlled Trials Register), and Best Evidence (American College of Physicians-American Society of Internal Medicine). Searches were limited to the English language, 1995 through 2000, and human subjects. In addition, related-article searches were conducted in MEDLINE to find further relevant articles. Finally, committee members recommended applicable articles outside the scope of the formal searches.

Major search topics included perioperative risk, cardiac risk, noncardiac surgery, noncardiac, intraoperative risk, postoperative risk, risk stratification, cardiac

complication, cardiac evaluation, perioperative care, preoperative evaluation, preoperative assessment, and intraoperative complications. Additional searches cross-referenced these topics with the following subtopics: troponin, myocardial infarction, myocardial ischemia, Duke activity status index, functional capacity, dobutamine, adenosine, venous thrombosis, thromboembolism, warfarin, percutaneous transluminal coronary angioplasty (PTCA), adrenergic beta-agonists, echocardiography, anticoagulant, beta-blocker, diabetes mellitus, wound infection, blood sugar control, normothermia, body temperature changes, body temperature regulation, hypertension, pulmonary hypertension, anemia, aspirin, arrhythmia, implantable defibrillator, artificial pacemaker, pulmonary artery catheters, Swan Ganz catheter, and platelet aggregation inhibitors.

NUMBER OF SOURCE DOCUMENTS

400 relevant, new articles were identified and reviewed by the committee for the update of the guideline.

METHODS USED TO ASSESS THE QUALITY AND STRENGTH OF THE EVIDENCE

Weighting According to a Rating Scheme (Scheme Not Given)

RATING SCHEME FOR THE STRENGTH OF THE EVIDENCE

Highest priority given to randomized trials; second highest priority given to observational database reports; lowest priority given to expert opinion

METHODS USED TO ANALYZE THE EVIDENCE

Systematic Review with Evidence Tables

DESCRIPTION OF THE METHODS USED TO ANALYZE THE EVIDENCE

The committee conducted a systematic review with thorough evaluation of available clinical data and compilation of these data into six evidence tables.

METHODS USED TO FORMULATE THE RECOMMENDATIONS

Expert Consensus

DESCRIPTION OF METHODS USED TO FORMULATE THE RECOMMENDATIONS

Experts in the subject under consideration are selected from the American College of Cardiology and the American Heart Association to examine subject-specific data and write guidelines. The process includes additional representatives from other medical specialty groups when appropriate. Writing groups are specifically charged to perform a formal literature review, weigh the strength of evidence for or against a particular treatment or procedure, and include estimates of expected health outcomes where data exist. Patient-specific modifiers, comorbidities, and

issues of patient preference that might influence the choice of particular tests or therapies are considered as well as frequency of follow-up and cost-effectiveness.

RATING SCHEME FOR THE STRENGTH OF THE RECOMMENDATIONS

Class I: Conditions for which there is evidence for and/or general agreement that the procedure or treatment is useful and effective.

Class II: Conditions for which there is conflicting evidence and/or a divergence of opinion about the usefulness/ efficacy of a procedure or treatment.

Class II a: The weight of evidence or opinion is in favor of the procedure or treatment.

Class IIb: Usefulness/efficacy is less well established by evidence or opinion.

Class III: Conditions for which there is evidence and/or general agreement that the procedure or treatment is not useful/effective and in some cases may be harmful.

COST ANALYSIS

Implications of Risk Assessment Strategies for Costs

The decision to recommend further noninvasive or invasive testing for the individual patient being considered for noncardiac surgery ultimately becomes a balancing act between the estimated probabilities of effectiveness vs. risk. The proposed benefit, of course, is the possibility of identifying advanced but relatively unsuspected coronary artery disease (CAD) that might result in significant cardiac morbidity or mortality either perioperatively or in the long term. In the process of further screening and treatment, the risks from the tests and treatments themselves may offset or even exceed the potential benefit of evaluation. Furthermore, the cost of screening and treatment strategies must be considered. Although physicians should be concerned with improving the clinical outcome of their patients, cost is an appropriate consideration when different evaluation and treatment strategies are available that cannot be distinguished from one another in terms of clinical outcome.

Formal decision and cost-effectiveness analyses of this particular question have been done and have yielded highly varied results. Because the exact amount of risk reduction from coronary revascularization in the clinical populations differs so much from center to center, it is difficult to determine the exact risks of aggressive screening and treatments vs. the benefits in terms of risk reduction. Additionally, the models all demonstrate that optimal strategy depends on the mortality rates for both cardiac procedures and noncardiac surgeries in the clinically relevant range. One decision model, which did not support a strategy incorporating coronary angiography and revascularization, used lower mortality rates than those used or reported in the other studies. Therefore, use of any decision and cost-effectiveness model in a specific situation depends on the comparability of local mortality rates to those of the model.

One report suggested that the cost of a selected coronary screening approach, as described in these guidelines, was as low as \$214 per patient. Several recent publications have shown a cost per year of life saved for this selected screening strategy of less than \$45,000 when applied to patients undergoing vascular surgery. However, none of these studies included a strategy of selected screening followed by aggressive beta-blocker treatment in high-risk individuals, as recently described by Poldermans and colleagues. It is likely that this approach will be preferred over more aggressive coronary assessment/treatment strategies except perhaps among very high-risk subsets of patients. Prophylactic beta-blockade represents an excellent strategy in patients for whom coronary revascularization for long-term benefit is not a serious consideration.

Postoperative and Long-Term Management

In general, the indications for additional screening or testing in postoperative patients depend on individual patient characteristics. A recent decision-tree model was constructed to compare cost-effectiveness of various preoperative screening protocols in postoperative vascular surgery patients for up to 5 years after discharge. The best event-free survival and cost-effectiveness ratio were noted for selective preoperative stress testing (using dipyridamole-thallium imaging) in patients with intermediate clinical risk, whereas high-risk patients were referred to coronary angiography and low-risk patients were sent to elective surgery without further workup. This is the general approach suggested in these guidelines.

METHOD OF GUIDELINE VALIDATION

External Peer Review Internal Peer Review

DESCRIPTION OF METHOD OF GUIDELINE VALIDATION

This update was reviewed by two outside reviewers from the American Heart Association and two outside reviewers of the American College of Cardiology, as well as one reviewer of the American College of Cardiology/American Heart Association Task Force on Practice Guidelines. It was approved by the American College or Cardiology Board of Trustees and the American Heart Association Science Advisory and Coordinating Committee.

RECOMMENDATIONS

MAJOR RECOMMENDATIONS

Excerpted by the National Guideline Clearinghouse (NGC)

Explanation of classification system for the guideline recommendations is provided at the end of the "Major Recommendations" field.

General Approach

The preoperative cardiac evaluation must be carefully tailored to the circumstances that have prompted the consultation and to the nature of the

surgical illness (e.g., acute surgical emergency) as opposed to urgent or elective cases. Successful perioperative evaluation and treatment of cardiac patients undergoing noncardiac surgery requires careful teamwork and communication between the patient, primary care physician, anesthesiologist, consultant, and surgeon. In general, indications for further cardiac testing and treatments are the same as those in the nonoperative setting, but their timing is dependent on such factors as the urgency of noncardiac surgery, the patient 's risk factors, and specific surgical considerations. Coronary revascularization before noncardiac surgery to enable the patient to "get through" the noncardiac procedure is appropriate only for a small subset of patients at very high risk. Preoperative testing should be limited to circumstances in which the results will affect patient treatment and outcomes. A conservative approach to the use of expensive tests and treatments is recommended.

Preoperative Clinical Evaluation

The initial history, physical examination, and electrocardiogram (ECG) assessment should focus on identification of potentially serious cardiac disorders, including coronary artery disease (CAD) [e.g., prior myocardial infarction (MI) and angina pectoris], heart failure (HF), symptomatic arrhythmias, presence of pacemaker or implantable cardioverter defibrillator (ICD), or a history of orthostatic intolerance. The presence of anemia may also place a patient at higher perioperative risk.

In addition to identifying the presence of preexisting manifested heart disease, it is essential to define disease severity, stability, and prior treatment. Other factors that help determine cardiac risk include functional capacity, age, comorbid conditions (e.g., diabetes mellitus, peripheral vascular disease, renal dysfunction, and chronic pulmonary disease), and type of surgery (vascular procedures and prolonged, complicated thoracic, abdominal, and head and neck procedures are considered higher risk).

Numerous risk indices have been developed over the past 25 years on the basis of multivariate analyses. In addition to the presence of CAD and heart failure, a history of cerebrovascular disease, preoperative elevated creatinine greater than 2 mg per deciliter, insulin treatment for diabetes mellitus, and high-risk surgery have all been associated with increased perioperative cardiac morbidity. Despite these risk indices, there was consensus among the committee members to place clinical risk factors into 3 categories of predictors (see the section titled "Clinical Markers," below).

Further Preoperative Testing to Assess Coronary Risk

Which patients are most likely to benefit from preoperative coronary assessment and treatment? The lack of adequately controlled or randomized clinical trials to define the optimal evaluation strategy led to the proposed algorithm based on collected observational data and expert opinion (see Figure 1, <u>Stepwise Approach to Preoperative Cardiac Assessment</u>, in the original guideline document). Since publication of the guidelines in 1996, several studies have suggested that this stepwise approach to the assessment of CAD is both efficacious and cost-effective.

A stepwise Bayesian strategy that relies on assessment of clinical markers, prior coronary evaluation and treatment, functional capacity, and surgery-specific risk

is outlined in Figure 1 in the original guideline document. A framework for determining which patients are candidates for cardiac testing is presented in algorithmic form (see Figure 1, <u>Stepwise Approach to Preoperative Cardiac Assessment</u>, in the original guideline document). Successful use of the algorithm requires an appreciation of the different levels of risk attributable to certain clinical circumstances, levels of functional capacity, and types of surgery. These are defined below, after which the algorithm is reviewed step by step.

Clinical Markers

The major clinical predictors (see Table 1, below) of increased perioperative cardiovascular risk are a recent unstable coronary syndrome such as an acute MI (documented MI less than 7 days previously), recent MI (more than 7 days but less than 1 month before surgery), unstable or severe angina, evidence of a large ischemic burden by clinical symptoms or noninvasive testing, decompensated heart failure, significant arrhythmias (high-grade atrioventricular block, symptomatic arrhythmias in the presence of underlying heart disease, or supraventricular arrhythmias with uncontrolled ventricular rate), and severe valvular disease.

Intermediate predictors of increased risk are mild angina pectoris, a more remote prior MI (more than 1 month before planned surgery), compensated heart failure, preoperative creatinine greater than or equal to 2.0 mg per deciliter, and diabetes mellitus. Minor predictors of risk are advanced age, abnormal ECG, rhythm other than sinus, low functional capacity, history of stroke, and uncontrolled systemic hypertension.

A history of MI or abnormal Q waves by ECG is listed as an intermediate predictor, whereas an acute MI (defined as at least 1 documented MI less than or equal to 7 days before the examination) or recent MI (more than 7 days but less than or equal to 1 month before the examination) with evidence of important ischemic risk by clinical symptoms or noninvasive study is a major predictor. This definition reflects the consensus of the American College of Cardiology Cardiovascular Database Committee. In this way, the separation of MI into the traditional 3- and 6-month intervals has been avoided. Current management of MI provides for risk stratification during convalescence. If a recent stress test does not indicate residual myocardium at risk, the likelihood of reinfarction after noncardiac surgery is low. Although there are no adequate clinical trials on which to base firm recommendations, it appears reasonable to wait 4 to 6 weeks after MI to perform elective surgery.

Table 1. Clinical Predictors of Increased Perioperative Cardiovascular Risk (Myocardial Infarction, Heart Failure, Death)

Major

- Unstable coronary syndromes
 - Acute or recent myocardial infarction* with evidence of important ischemic risk by clinical symptoms or noninvasive study

- Unstable or severe** angina (Canadian class III or IV)***
- Decompensated heart failure
- Significant arrhythmias
 - High-grade atrioventricular block
 - Symptomatic ventricular arrhythmias in the presence of underlying heart disease
 - Supraventricular arrhythmias with uncontrolled ventricular rate
- Severe valvular disease

Intermediate

- Mild angina pectoris (Canadian class I or II)
- Previous myocardial infarction by history or pathological Q waves
- Compensated or prior heart failure
- Diabetes mellitus (particularly insulin-dependent)
- Renal insufficiency

Minor

- Advanced age
- Abnormal ECG (left ventricular hypertrophy, left bundle-branch block, ST-T abnormalities)
- Rhythm other than sinus (e.g., atrial fibrillation)
- Low functional capacity (e.g., inability to climb one flight of stairs with a bag of groceries)
- History of stroke
- Uncontrolled systemic hypertension
- *The American College of Cardiology National Database Library defines recent myocardial infarction (MI) as greater than 7 days but less than or equal to 1 month (30 days); acute myocardial infarction is within 7 days.
- **May include "stable" angina in patients who are unusually sedentary.
- ***Campeau L. Grading of angina pectoris. Circulation 1976;54:522–3.

Functional Capacity

Functional capacity can be expressed in metabolic equivalent (MET) levels (see Table 2, below). Multiples of the baseline metabolic equivalent value can be used to express aerobic demands for specific activities. Perioperative cardiac and long-term risks are increased in patients unable to meet a 4-metabolic equivalent demand during most normal daily activities. The Duke Activity Status Index and other activity scales provide the clinician with a set of questions to determine a patient 's functional capacity. Energy expenditures for activities such as eating, dressing, walking around the house, and dishwashing range from 1 to 4 metabolic equivalents. Climbing a flight of stairs, walking on level ground at 6.4 km per hour, running a short distance, scrubbing floors, or playing a game of golf

represents 4 to 10 metabolic equivalents. Strenuous sports such as swimming, singles tennis, and football often exceed 10 metabolic equivalents.

Table 2. Estimated Energy Requirements for Various Activities*

1 metabolic equivalent. Can you take care of yourself? Eat, dress, or use the toilet? Walk indoors around the house? Walk a block or two on level ground at 2 to 3 mph or 3.2 to 4.8 km per h?

4 metabolic equivalents. Do light work around the house like dusting or washing dishes? Climb a flight of stairs or walk up a hill? Walk on level ground at 4 mph or 6.4 km per h? Run a short distance? Do heavy work around the house like scrubbing floors or lifting or moving heavy furniture? Participate in moderate recreational activities like golf, bowling, dancing, doubles tennis, or throwing a baseball or football?

Greater than 10 metabolic equivalents. Participate in strenuous ports like swimming, singles tennis, football, basketball, or skiing?

*Adapted from the Duke Activity Status Index (Hlatky MA, Boineau RE, Higginbotham MB, et al. A brief self-administered questionnaire to determine functional capacity [the Duke Activity Status Index]. Am J Cardiol 1989; 64: 651-4) and American Heart Association Exercise Standards (Fletcher GF, Balady G, Froelicher VF, Hartley LH, Haskell WL, Pollock ML. Exercise standards: a statement for healthcare professionals from the American Heart Association: Writing Group. Circulation 1995; 91: 580-615).

Surgery-Specific Risk

Surgery-specific cardiac risk of noncardiac surgery is related to 2 important factors: the type of surgery itself and the degree of hemodynamic stress associated with the procedures. The duration and intensity of coronary and myocardial stressors can be helpful in estimating the likelihood of perioperative cardiac events, particularly for emergency surgery. Surgery-specific risk for noncardiac surgery can be stratified as high, intermediate, and low (see Table 3, below).

Table 3. Cardiac Risk* Stratification for Noncardiac Surgical Procedures

- High (Reported cardiac risk often greater than 5%)
 - Emergent major operations, particularly in the elderly
 - Aortic and other major vascular surgery
 - Peripheral vascular surgery
 - Anticipated prolonged surgical procedures associated with large fluid shifts and/or blood loss
- Intermediate (Reported cardiac risk generally less than 5%)
 - Carotid endarterectomy

- Head and neck surgery
- Intraperitoneal and intrathoracic surgery
- Orthopedic surgery
- Prostate surgery
- Low** (Reported cardiac risk generally less than 1%)
 - Endoscopic procedures
 - Superficial procedure
 - Cataract surgery
 - Breast surgery
- *Combined incidence of cardiac death and nonfatal myocardial infarction.
- **Do not generally require further preoperative cardiac testing.

An algorithm for the <u>Stepwise Approach to Preoperative Cardiac Assessment</u> is available from the American College of Cardiology Web site. The following steps correspond to that algorithm:

- Step 1. What is the urgency of noncardiac surgery? Certain emergencies do not allow time for preoperative cardiac evaluation. Postoperative risk stratification may be appropriate for some patients who have not had such an assessment before.
- Step 2. Has the patient undergone coronary revascularization in the past 5 years? If so, and if clinical status has remained stable without recurrent symptoms/signs of ischemia, further cardiac testing is generally not necessary.
- Step 3. Has the patient had a coronary evaluation in the past 2 years? If coronary risk was adequately assessed and the findings were favorable, it is usually not necessary to repeat testing unless the patient has experienced a change or new symptoms of coronary ischemia since the previous evaluation.
- Step 4. Does the patient have an unstable coronary syndrome or a major clinical predictor of risk? When elective noncardiac surgery is being considered, the presence of unstable coronary disease, decompensated heart failure, symptomatic arrhythmias, and/or severe valvular heart disease usually leads to cancellation or delay of surgery until the problem has been identified and treated.
- Step 5. Does the patient have intermediate clinical predictors of risk? The presence or absence of prior MI by history or ECG, angina pectoris, compensated or prior heart failure, preoperative creatinine greater than or equal to 2 mg per deciliter, and/or diabetes mellitus helps to further stratify clinical risk for perioperative coronary events. Consideration of functional capacity and level of surgery-specific risk allows a rational approach to identify patients most likely to benefit from further noninvasive testing.
- Step 6. Patients without major but with intermediate predictors of clinical risk and moderate or excellent functional capacity can generally undergo intermediate-risk surgery with little likelihood of perioperative death or MI. Conversely, further noninvasive testing is often considered for patients with poor functional capacity or moderate functional capacity but higher-risk surgery, especially for patients with 2 or more intermediate predictors of risk.

Step 7. Noncardiac surgery is generally safe for patients with neither major nor intermediate predictors of clinical risk and moderate or excellent functional capacity (4 metabolic equivalents or greater). Additional testing may be considered on an individual basis for patients without clinical markers but with poor functional capacity who are facing higher-risk operations, particularly those with several minor clinical predictors of risk who are scheduled to undergo vascular surgery.

Step 8. The results of noninvasive testing can be used to determine the need for additional preoperative testing and treatment. In some patients with documented CAD, the risk of coronary intervention or corrective cardiac surgery may approach or even exceed the risk of the proposed noncardiac surgery. This approach may be appropriate, however, if it significantly improves the patient 's long-term prognosis.

For some patients, a careful consideration of clinical, surgery-specific, and functional status attributes leads to a decision to proceed to coronary angiography.

Management of Specific Preoperative Cardiovascular Conditions

Hypertension

Stage 3 hypertension (systolic blood pressure greater than or equal to 180 mmHg and diastolic blood pressure greater than or equal to 110 mmHg) should be controlled before surgery. In many such instances, establishment of an effective regimen can be achieved over several days to weeks of preoperative outpatient treatment. If surgery is more urgent, rapid-acting agents can be administered that allow effective control in a matter of minutes or hours. Beta-blockers appear to be particularly attractive agents. Continuation of preoperative antihypertensive treatment through the perioperative period is critical.

Valvular Heart Disease

Indications for evaluation and treatment of valvular heart disease are identical to those in the nonpreoperative setting. Symptomatic stenotic lesions are associated with risk of perioperative heart failure or shock and often require percutaneous valvotomy or valve replacement before noncardiac surgery to lower cardiac risk. Symptomatic regurgitant valve disease is usually better tolerated perioperatively and may be stabilized preoperatively with intensive medical therapy and monitoring. Regurgitant valve disease can then be treated definitively with valve repair or replacement after noncardiac surgery. Medical therapy and monitoring are appropriate when a delay of several weeks or months before noncardiac surgery may have severe consequences. Exceptions may include severe valvular regurgitation with reduced left ventricular function, in which overall hemodynamic reserve is so limited that destabilization during perioperative stresses is likely.

Myocardial Disease

Dilated and hypertrophic cardiomyopathy are associated with an increased incidence of perioperative heart failure. Management is aimed at maximizing

preoperative hemodynamic status and providing intensive postoperative medical therapy and surveillance. An estimate of hemodynamic reserve is useful for anticipating potential complications from intraoperative or postoperative stress.

Arrhythmias and Conduction Abnormalities

The presence of an arrhythmia or cardiac conduction disturbance should provoke a careful evaluation for under-lying cardiopulmonary disease, drug toxicity, or metabolic abnormality. Therapy should be initiated for symptomatic or hemodynamically significant arrhythmias, first to reverse an underlying cause and second to treat the arrhythmia. Indications for antiarrhythmic therapy and cardiac pacing are identical to those in the nonoperative setting. Frequent ventricular premature beats and/or asymptomatic nonsustained ventricular tachycardia have not been associated with an increased risk of nonfatal MI or cardiac death in the perioperative period, and therefore, aggressive monitoring or treatment in the perioperative period generally is not necessary.

Implantable Pacemakers or Implantable Cardioverter Defibrillators (ICDs)

The type and extent of evaluation of a pacemaker or implantable cardioverter defibrillator depend on the urgency of the surgery, whether a pacemaker has unipolar or bipolar leads, whether electrocautery is bipolar or unipolar, the distance between electrocautery and pacemaker, and pacemaker dependency. implantable cardioverter defibrillator devices should be programmed off immediately before surgery and then on again postoperatively.

Supplemental Preoperative Evaluation

Specific recommendations for supplemental preoperative evaluation must be individualized to each patient and circumstance. The following may be appropriate in specific situations: assessment of resting left ventricular function, exercise stress testing, pharmacological stress testing, ambulatory ECG monitoring, and coronary angiography. In most ambulatory patients, the test of choice is exercise ECG testing, which can both provide an estimate of functional capacity and detect myocardial ischemia through changes in the ECG and hemodynamic response. In patients with important abnormalities on their resting ECG (e.g., left bundle-branch block, left ventricular hypertrophy with strain pattern, or digitalis effect), other techniques such as exercise echocardiography or exercise myocardial perfusion imaging should be considered. Recommendations regarding individual testing modalities are given below.

Recommendations for Preoperative Noninvasive Evaluation of Left Ventricular Function

Class I. Patients with current or poorly controlled heart failure. (If previous evaluation has documented severe left ventricular dysfunction, repeat preoperative testing may not be necessary.)

Class II a. Patients with prior heart failure and patients with dyspnea of unknown origin.

Class III. As a routine test of left ventricular function in patients without prior heart failure.

Recommendations for Preoperative 12-Lead Rest ECG

Class I. Recent episode of chest pain or ischemic equivalent in clinically intermediate- or high-risk patients scheduled for an intermediate- or high-risk operative procedure.

Class IIa. Asymptomatic persons with diabetes mellitus.

Class IIb.

- 1. Patients with prior coronary revascularization.
- 2. Asymptomatic male more than 45 years old or female more than 55 years old with 2 or more atherosclerotic risk factors.
- 3. Prior hospital admission for cardiac causes.

Class III. As a routine test in asymptomatic subjects undergoing low-risk operative procedures.

Recommendations for Exercise or Pharmacological Stress Testing

Class I.

- 1. Diagnosis of adult patients with intermediate pretest probability of CAD.
- 2. Prognostic assessment of patients undergoing initial evaluation for suspected or proven CAD; evaluation of subjects with significant change in clinical status.
- 3. Demonstration of proof of myocardial ischemia before coronary revascularization.
- 4. Evaluation of adequacy of medical therapy; prognostic assessment after an acute coronary syndrome (if recent evaluation unavailable).

Class II a. Evaluation of exercise capacity when subjective assessment is unreliable.

Class IIb.

- 1. Diagnosis of CAD patients with high or low pretest probability: those with resting ST depression less than 1 mm, those undergoing digitalis therapy, or those with ECG criteria for left ventricular hypertrophy.
- 2. Detection of restenosis in high-risk asymptomatic subjects within the initial months after percutaneous coronary intervention (PCI).

Class III.

1. For exercise stress testing, diagnosis of patients with resting ECG abnormalities that preclude adequate assessment, e.g., pre-excitation syndrome, electronically paced ventricular rhythm, rest ST depression greater than 1 mm, or left bundle-branch block.

- Severe comorbidity likely to limit life expectancy or candidacy for revascularization.
- 3. Routine screening of asymptomatic men or women without evidence of CAD.
- 4. Investigation of isolated ectopic beats in young patients.

Recommendations for Coronary Angiography in Perioperative Evaluation Before (or After) Noncardiac Surgery

Class I. Patients With Suspected or Known CAD

- 1. Evidence for high risk of adverse outcome based on noninvasive test results.
- 2. Angina unresponsive to adequate medical therapy.
- 3. Unstable angina, particularly when facing intermediate-risk* or high-risk* noncardiac surgery.
- 4. Equivocal noninvasive test results in patients at high clinical risk** undergoing high-risk* surgery.

Class IIa.

- 1. Multiple markers of intermediate clinical risk** and planned vascular surgery (noninvasive testing should be considered first).
- 2. Moderate to large region of ischemia on noninvasive testing but without highrisk features and lower left ventricular ejection fraction.
- 3. Nondiagnostic noninvasive test results in patients of intermediate clinical risk** undergoing high-risk* noncardiac surgery.
- 4. Urgent noncardiac surgery while convalescing from acute myocardial infarction.

Class IIb.

- 1. Perioperative MI.
- Medically stabilized class III or IV angina and planned low-risk or minor* surgery.

Class III.

- 1. Low-risk* noncardiac surgery with known coronary artery disease (CAD) and no high-risk results on noninvasive testing.
- 2. Asymptomatic after coronary revascularization with excellent exercise capacity (greater than or equal to 7 metabolic equivalents [METs]).
- 3. Mild stable angina with good left ventricular function and no high-risk noninvasive test results.
- 4. Noncandidate for coronary revascularization owing to concomitant medical illness, severe left ventricular dysfunction (e.g., left ventricular ejection fraction less than 0.20), or refusal to consider revascularization.
- 5. Candidate for liver, lung, or renal transplant less than 40 years old, as part of evaluation for transplantation, unless noninvasive testing reveals high risk for adverse outcome.

^{*}Cardiac risk according to type of noncardiac surgery. High risk: emergent major operations, aortic and major vascular, peripheral vascular, or anticipated

prolonged surgical procedure associated with large fluid shifts and blood loss; intermediate risk: carotid endarterectomy, major head and neck, intraperitoneal and intrathoracic, orthopedic, prostate; and low risk: endoscopic procedures, superficial procedures, cataract, or breast.

**Cardiac risk according to clinical predictors of perioperative death, MI, or heart failure. High clinical risk: unstable angina, acute or recent MI, and evidence of important residual ischemic risk, decompensated heart failure, high degree of atrioventricular block, symptomatic ventricular arrhythmias with known structural heart disease, severe symptomatic valvular heart disease, patient with multiple intermediate-risk markers such as prior MI, heart failure, and diabetes; intermediate clinical risk: Canadian Cardiovascular Society class I or II angina, prior MI by history or ECG, compensated or prior heart failure, diabetes mellitus or renal insufficiency.

Perioperative Therapy or Previous Coronary Revascularization

Coronary Artery Bypass Grafting

Indications for coronary artery bypass grafting (CABG) before noncardiac surgery are identical to those reviewed in the American College of Cardiology/American Heart Association guidelines for coronary artery bypass grafting (see the <u>National Guideline Clearinghouse Guideline Summary</u>). Coronary artery bypass grafting is rarely indicated simply to "get a patient through" noncardiac surgery. Patients undergoing elective noncardiac procedures who are found to have prognostic high-risk coronary anatomy and in whom long-term outcome would likely be improved by coronary artery bypass grafting should generally undergo revascularization before a noncardiac elective surgical procedure of high or intermediate risk (see Table 3, above).

Percutaneous Coronary Intervention

Until further data are available, indications for percutaneous coronary intervention in the perioperative setting are similar to those in the American College of Cardiology/American Heart Association guidelines for use of percutaneous coronary intervention in general (see the <u>National Guideline Clearinghouse Guideline Summary</u>). There is uncertainty regarding how much time should pass between percutaneous coronary intervention and noncardiac procedures. Delaying surgery for at least 1 week after balloon angioplasty to allow for healing of the vessel injury has theoretical benefits. If a coronary stent is used, a delay of at least 2 weeks and ideally 4 to 6 weeks should occur before noncardiac surgery to allow 4 full weeks of dual antiplatelet therapy and re-endothelialization of the stent to be completed, or nearly so.

Perioperative Medical Therapy

Recommendations for Perioperative Medical Therapy

Class L.

- 1. Beta-blockers required in the recent past to control symptoms of angina or patients with symptomatic arrhythmias or hypertension.
- 2. Beta-blockers: patients at high cardiac risk owing to the finding of ischemia on preoperative testing who are undergoing vascular surgery.

Class IIa.

1. Beta-blockers: preoperative assessment identifies untreated hypertension, known coronary disease, or major risk factors for coronary disease.

Class IIb.

1. Alpha-2 agonist: perioperative control of hypertension, or known CAD or major risk factors for CAD.

Class III.

- 1. Beta-blockers: contraindication to beta-blockade.
- 2. Alpha-2 agonists: contraindication to alpha-2 agonists.

Anesthetic Considerations and Intraoperative Management

Anesthetic Agent

All anesthetic techniques and drugs have known cardiac effects that should be considered in the perioperative plan. There appears to be no one best myocardium-protective anesthetic technique. Therefore, the choice of anesthesia and intraoperative monitors is best left to the discretion of the anesthesia care team, which will consider the need for postoperative ventilation, cardiovascular effects (including myocardial depression), sympathetic blockade, and dermatomal level of the procedure. Advocates of monitored anesthesia, in which local anesthesia is supplemented by intravenous sedation/analgesia, have argued that use of this technique avoids the undesirable effects of general or neuraxial techniques, but no studies have established this. Failure to produce complete local anesthesia/analgesia can lead to increased stress response and/or myocardial ischemia.

Perioperative Pain Management

Patient-controlled intravenous and/or epidural analgesia is a popular method for reducing postoperative pain. Several studies suggest that effective pain management leads to a reduction in postoperative catecholamine surges and hypercoagulability.

Intraoperative Nitroglycerin

There are insufficient data about the effects of prophylactic intraoperative intravenous nitroglycerin in patients at high risk. Nitroglycerin should be used only when the hemodynamic effects of other agents in use have been considered.

Recommendations for Intraoperative Nitroglycerin

Class I. High-risk patients previously taking nitroglycerin who have active signs of myocardial ischemia without hypotension.

Class IIb. As a prophylactic agent for high-risk patients to prevent myocardial ischemia and cardiac morbidity, particularly in those who have required nitrate therapy to control angina. The recommendation for prophylactic use of nitroglycerin must take into account the anesthetic plan and patient hemodynamics and must recognize that vasodilation and hypovolemia can readily occur during anesthesia and surgery.

Class III. Patients with signs of hypovolemia or hypotension.

Transesophageal Echocardiography

There are few data on the value of transesophageal echocardiography to detect transient wall motion abnormalities in predicting cardiac morbidity in noncardiac surgical patients. Experience to date suggests that the incremental value of this technique for risk prediction is small. Guidelines for appropriate use of transesophageal echocardiography have been published by the American Society of Anesthesiologists and the Society of Cardiovascular Anesthesiologists (Practice guidelines for perioperative transesophageal echocardiography: a report by the American Society of Anesthesiologists and the Society of Cardiovascular Anesthesiologists Task Force on Transesophageal Echocardiography. Anesthesiology 1996;84: 986-1006).

Perioperative Maintenance of Body Temperature

One randomized trial demonstrated a reduced incidence of perioperative cardiac events in patients who were maintained in a state of normothermia via forced-air warming compared with routine care.

Perioperative Surveillance

Pulmonary Artery Catheters

Although very few studies that have been reported compare patient outcomes after treatment with or without pulmonary artery catheters, 3 variables are particularly important in assessing benefit versus risk of pulmonary artery catheter use: disease severity, magnitude of anticipated surgery, and practice setting. The extent of expected fluid shifts is a primary concern. Patients most likely to benefit from perioperative use of a pulmonary artery catheter appear to be those with a recent myocardial infarction complicated by heart failure, those with significant CAD who are undergoing procedures associated with significant hemodynamic stress, and those with systolic or diastolic left ventricular dysfunction, cardiomyopathy, and/or valvular disease who are undergoing highrisk operations.

Intraoperative and Postoperative ST-Segment Monitoring

Intraoperative and postoperative ST changes indicating myocardial ischemia are strong predictors of perioperative myocardial infarction in patients at high risk

who undergo noncardiac surgery. Similarly, postoperative ischemia is a significant predictor of long-term risk of myocardial infarction and cardiac death. Conversely, in patients at low risk who undergo noncardiac surgery, ST depression may occur and often is not associated with regional wall-motion abnormalities. Accumulating evidence suggests that proper use of computerized ST-segment analysis in appropriately selected patients at high risk may improve sensitivity for myocardial ischemia detection.

Recommendations for Perioperative ST-Segment Monitoring

Class II a. When available, proper use of computerized ST-segment analysis in patients with known CAD or undergoing vascular surgery may provide increased sensitivity to detect myocardial ischemia during the perioperative period and may identify patients who would benefit from further postoperative and long-term interventions.

Class IIb. Patients with single or multiple risk factors for CAD.

Class LLL Patients at low risk for CAD.

Surveillance for Perioperative MI

Few studies have examined the optimal method for diagnosing a perioperative MI. Clinical symptoms, postoperative ECG changes, and elevation of the MB fraction of creatine kinase (CK-MB) have been studied most extensively. Recently, elevations of myocardium-specific enzymes such as troponin-I, troponin-T, or CK-MB isoforms have also been shown to be of value (85 to 90). In patients with known or suspected CAD who are undergoing high-risk procedures, ECGs obtained at baseline, immediately after surgery, and on the first 2 days after surgery appear to be cost-effective. A risk gradient can be based on the magnitude of biomarker elevation, the presence or absence of concomitant new ECG abnormalities, hemodynamic instability, and quality and intensity of chest pain syndrome, if present. Use of cardiac biomarkers is best reserved for patients at high risk and those with clinical, ECG, or hemodynamic evidence of cardiovascular dysfunction.

Postoperative and Long-Term Management

Despite even optimal perioperative management, some patients will have perioperative MI, which is associated with a 40% to 70% mortality rate. For patients who experience a symptomatic perioperative ST-segment elevation MI as a result of sudden thrombotic coronary occlusion, angioplasty should be considered after the risks versus benefits have been weighed. Pharmacological therapy with aspirin should be initiated as soon as possible, and a beta-blocker and angiotensin converting enzyme inhibitor may also be beneficial. Perioperative MI carries a high risk for future cardiac events. Patients who sustain acute MI in the perioperative period should receive careful medical evaluation for residual ischemia and overall left ventricular function.

It is also appropriate to recommend secondary risk reduction in the relatively large number of elective surgery patients in whom cardiovascular abnormalities are detected during preoperative evaluations. Although the occasion of surgery is

often taken as a specific high-risk time, most of the patients who have known or newly detected CAD during their preoperative evaluations will not have any events during elective noncardiac surgery. After the preoperative cardiac risk has been determined by clinical or noninvasive testing, most patients will benefit from pharmacological agents to lower low-density lipoprotein cholesterol levels, increase high-density lipoprotein levels, or both. On the basis of expert opinion, the goal should be to lower the low-density lipoprotein level to less than 100 mg per deciliter (2.6 mmol per deciliter).

<u>American College of Cardiology/American Heart Association</u> Classifications of Evidence:

Class I: Conditions for which there is evidence for and/or general agreement that the procedure/therapy is useful and effective

Class II: Conditions for which there is conflicting evidence and/or a divergence of opinion about the usefulness/efficacy of performing the procedure/therapy

Class IIa: Weight of evidence/opinion is in favor of usefulness/efficacy.

Class IIb: Usefulness/efficacy is less well established by evidence/opinion.

Class III: Conditions for which there is evidence and/or general agreement that the procedure/therapy is not useful/effective and in some cases may be harmful.

CLINICAL ALGORITHM(S)

Clinical algorithms are provided for the <u>Stepwise Approach to Preoperative Cardiac Assessment</u> and for choosing an <u>Appropriate preoperative stress test</u>.

EVIDENCE SUPPORTING THE RECOMMENDATIONS

TYPE OF EVIDENCE SUPPORTING THE RECOMMENDATIONS

The type of evidence is not specifically stated for each recommendation. Observational or retrospective data and expert opinion form the basis of the proposed algorithm for preoperative cardiac assessment. Data from observational studies for recommendations is compiled in six evidence tables in the original guideline document (see Tables 6 through 11).

BENEFITS/HARMS OF IMPLEMENTING THE GUIDELINE RECOMMENDATIONS

POTENTIAL BENEFITS

- Appropriate utilization of invasive and noninvasive tests to evaluate cardiac risk in patients undergoing noncardiac surgery
- Decreased perioperative risk and cardiovascular morbidity (e.g., myocardial infarction) and mortality

Not stated

QUALIFYING STATEMENTS

OUALIFYING STATEMENTS

- These guidelines attempt to define practices that meet the needs of most
 patients in most circumstances. The ultimate judgment regarding the care of
 a particular patient must be made by the physician and patient in light of all
 the circumstances presented by that patient.
- While the collective knowledge surrounding the identification of high-and low-risk patients using perioperative clinical and noninvasive evaluation is substantial, very few prospective or randomized studies have been performed that establish the value of tests or treatments on perioperative outcome. Because the studies were rarely randomized controlled trial, definitions of a perioperative event varied, investigators were rarely blinded, and many inherent selection biases existed, the task force has chosen not to provide an aggregate synthesis of the data in the form of a point estimate or meta-analysis.

IMPLEMENTATION OF THE GUIDELINE

DESCRIPTION OF IMPLEMENTATION STRATEGY

An implementation strategy was not provided.

INSTITUTE OF MEDICINE (IOM) NATIONAL HEALTHCARE QUALITY REPORT CATEGORIES

IOM CARE NEED

Living with Illness Staying Healthy

IOM DOMAIN

Effectiveness

IDENTIFYING INFORMATION AND AVAILABILITY

BIBLIOGRAPHIC SOURCE(S)

American College of Cardiology Foundation (ACCF), American Heart Association (AHA). ACC/AHA guideline update on perioperative cardiovascular evaluation for noncardiac surgery. A report of the American College of Cardiology/American Heart Association Task Force on practice guidelines (Committee to Update the

1996 Guidelines). Bethesda (MD): American College of Cardiology Foundation; 2002. 58 p. [390 references]

ADAPTATION

Not applicable: The guideline was not adapted from another source.

DATE RELEASED

1996 Mar 15 (revised 2002)

GUI DELI NE DEVELOPER(S)

American College of Cardiology Foundation - Medical Specialty Society American Heart Association - Professional Association

SOURCE(S) OF FUNDING

The American College of Cardiology Foundation and the American Heart Association. No outside funding accepted.

GUI DELI NE COMMITTEE

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FINANCIAL DISCLOSURES/CONFLICTS OF INTEREST

The American College of Cardiology/American Heart Association (ACC/AHA) Task Force on Practice Guidelines makes every effort to avoid any actual or potential conflicts of interest that might arise as a result of an outside relationship or personal interest of a member of the writing panel. Specifically, all members of the writing panel are asked to provide disclosure statements of all such

^{*}Former Task Force member during the writing effort.

relationships that might be perceived as real or potential conflicts of interest. These statements are reviewed by the parent task force, reported orally to all members of the writing panel at the first meeting, and updated yearly and as change occur.

GUI DELI NE STATUS

This is the current release of the guideline. This guideline revises a previously issued version (Guidelines for perioperative cardiovascular evaluation for noncardiac surgery: report of the American College of Cardiology/American Heart Association Task Force on Practice Guidelines. J Am Coll Cardiol 1996 Jun; 27[4]: 910-48).

These guidelines will be reviewed annually after publication and considered current unless the American College of Cardiology/American Heart Association Task Force on Practice Guidelines publishes another update or full revision or withdraws the guidelines from publication.

GUIDELINE AVAILABILITY

Electronic copies: Available from the <u>American College of Cardiology (ACC) Web</u> <u>site</u>; electronic copies are also available in Portable Document Format (PDF) from the <u>American Heart Association (AHA) Web site</u>

Print copies: Available from Resource Center, American College of Cardiology, 9111 Old Georgetown Road, Bethesda, Maryland 20814-1699 and from the American Heart Association, Office of Scientific Affairs, 7272 Greenville Avenue, Dallas, TX 75231-4596.

AVAILABILITY OF COMPANION DOCUMENTS

The following is available:

 ACC/AHA guideline update on perioperative cardiovascular evaluation for noncardiac surgery – executive summary: a report of the American College of Cardiology/American Heart Association Task Force on Practice Guidelines (Committee to Update the 1996 Guidelines on Perioperative Cardiovascular Evaluation for Noncardiac Surgery). J Am Coll Cardiol 2002 Feb 6; 39(3):542-53; Circulation. 2002 Mar 12; 105(10):1257-67; Anesth Analg. 2002 May; 94(5):1052-64.

Electronic copies: Available from the American College of Cardiology Web site:

- HTML Format
- Portable Document Format (PDF)

Also available in Portable Document Format (PDF) from the <u>American Heart</u> Association Web site.

Print copies: Available from Resource Center, American College of Cardiology, 9111 Old Georgetown Road, Bethesda, Maryland 20814-1699 and from the

American Heart Association, Office of Scientific Affairs, 7272 Greenville Avenue, Dallas, TX 75231-4596.

PATIENT RESOURCES

None available

NGC STATUS

This summary was completed by ECRI on June 30, 1998. The information was verified by the guideline developer on December 1, 1998. This summary was updated by ECRI on April 30, 2002. The updated information was verified by the guideline developer on August 7, 2002.

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Date Modified: 4/12/2004



